

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

CORRECTED VERSION

(19) World Intellectual Property Organization International Bureau



(43) International Publication Date
18 March 2004 (18.03.2004)

PCT

(10) International Publication Number
WO 2004/022569 A1

(51) International Patent Classification⁷: C07F 9/117, C07C 59/13, 59/68, A61K 31/683, 31/185, A61P 35/00

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(21) International Application Number:
PCT/US2003/027607

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(22) International Filing Date:
3 September 2003 (03.09.2003)

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(25) Filing Language: English

(26) Publication Language: English

Published:

— with international search report

(30) Priority Data:
60/407,239 3 September 2002 (03.09.2002) US

(48) Date of publication of this corrected version:

10 June 2004

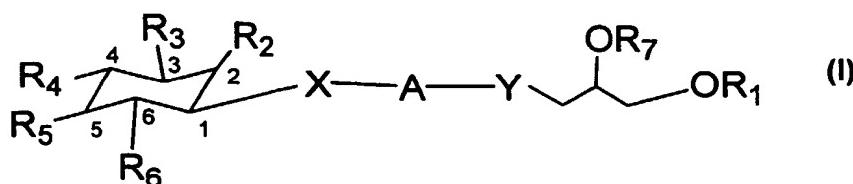
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(15) Information about Correction:

see PCT Gazette No. 24/2004 of 10 June 2004, Section II

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: AKT INHIBITORS, PHARMACEUTICAL COMPOSITIONS, AND USES THEREOF



inhibitors have the formula (I) wherein X and Y are independently selected from the group consisting of O, CF₃, CH₂, and CHF; wherein A is independently selected from the group consisting of P(O)OH, CH₂000H, and CH(COOH)₂; R₂ is selected from the group consisting of H, OH, isosteres of OH, C₁-C₂₅ alkoxy, C₆-C₁₀ aryloxy, C₃-C₈ cycloalkyloxy, C₃-C₈ cycloalkyl C₁-C₆ alkoxy, C₂-C₂₂ alkenyloxy, C₃-C₈ cycloalkenyloxy, C₇-C₃₂ aralkyloxy, C₇-C₃₂ alkylaryloxy, C₉-C₃₂ aralkenyl, and C₉-C₃₂ alkenylaryloxy; R₃-R₆ are independently selected from the group consisting of H, OH, isosteres of OH; and R₁ and R₇ are independently selected from the group consisting of C₁-C₂₅ alkyl, C₆-C₁₀ aryl, C₃-C₈ cycloalkyl, C₂-C₂₂ alkenyl, C₃-C₈ cycloalkenyl, C₇-C₃₂ aralkyl, C₇-C₃₂ alkylaryl, C₉-C₃₂ aralkenyl, and C₉-C₃₂ alkenylaryl; with the provisos that (i) when X is O, Y is O or CH₂, and R₃ is H, at least one of R₂ and R₄-R₆ is not OH; (ii) when A is CH₂COOH or CH(COOH)₂, X and Y cannot be simultaneously O; and (iii) all of R₂-R₆ are not simultaneously H. The inhibitors can be in the form of a salt also.

(57) Abstract: Disclosed are inhibitors of the serine/threonine kinase Akt, pharmaceutical compositions comprising such inhibitors, and a method of preventing or treating a disease or condition in an animal by the use of such inhibitors. The Akt

WO 2004/022569 A1